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TO

APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.
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09/509,712 03/31/00 RUBIN

D 01123.0004

HM22/0109

EXAMINER

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NEEDLE & ROSENBERG
127 PEACHTREE STREET NE
SUITE 1200 THE CANDLER BUILDING
ATLANTA GA 30303-1811

FOLEY,S

ART UNIT	PAPER NUMBER
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1648

5

DATE MAILED:

01/09/01

Please find below and/or attached an Office communication concerning this application or proceeding.

Commissioner of Patents and Trademarks

Office Action Summary	Application No.	Applicant(s)
	09/509,712	RUBIN ET AL.
	Examiner	Art Unit
	Shanon A. Foley	1648

-- The MAILING DATE of this communication appears on the cover sheet with the correspondence address --

Period for Reply

A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 1 MONTH(S) FROM THE MAILING DATE OF THIS COMMUNICATION.

- Extensions of time may be available under the provisions of 37 CFR 1.136 (a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.
- If the period for reply specified above is less than thirty (30) days, a reply within the statutory minimum of thirty (30) days will be considered timely.
- If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.
- Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133).
- Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).

Status

- 1) Responsive to communication(s) filed on ____ .
- 2a) This action is FINAL. 2b) This action is non-final.
- 3) Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under *Ex parte Quayle*, 1935 C.D. 11, 453 O.G. 213.

Disposition of Claims

- 4) Claim(s) 1-29 is/are pending in the application.
- 4a) Of the above claim(s) ____ is/are withdrawn from consideration.
- 5) Claim(s) ____ is/are allowed.
- 6) Claim(s) ____ is/are rejected.
- 7) Claim(s) ____ is/are objected to.
- 8) Claims 1-29 are subject to restriction and/or election requirement.

Application Papers

- 9) The specification is objected to by the Examiner.
- 10) The drawing(s) filed on ____ is/are objected to by the Examiner.
- 11) The proposed drawing correction filed on ____ is: a) approved b) disapproved.
- 12) The oath or declaration is objected to by the Examiner.

Priority under 35 U.S.C. § 119

- 13) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d).
- a) All b) Some * c) None of:
1. Certified copies of the priority documents have been received.
2. Certified copies of the priority documents have been received in Application No. ____ .
3. Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).
- * See the attached detailed Office action for a list of the certified copies not received.
- 14) Acknowledgement is made of a claim for domestic priority under 35 U.S.C. & 119(e).

Attachment(s)

- 15) Notice of References Cited (PTO-892)
- 16) Notice of Draftsperson's Patent Drawing Review (PTO-948)
- 17) Information Disclosure Statement(s) (PTO-1449) Paper No(s) ____ .
- 18) Interview Summary (PTO-413) Paper No(s) ____ .
- 19) Notice of Informal Patent Application (PTO-152)
- 20) Other: *Sequence Notice*

DETAILED ACTION

Election/Restriction

Restriction is required under 35 U.S.C. 121 and 372.

This application contains the following inventions or groups of inventions which are not so linked as to form a single general inventive concept under PCT Rule 13.1.

In accordance with 37 CFR 1.499, applicant is required, in response to this action, to elect a single invention to which the claims must be restricted.

Group I, claim(s) 1-14 and 20-23, drawn to a first product of nucleic acids. Election of this group requires an election of species.

Group II, claim(s) 15 and 16, drawn to a second product of nucleic acids comprising a regulatory region. Election of this group requires an election of species.

Group III, claim(s) 17-19, drawn to a method to inhibit viral infection. Election of this group requires an election of species.

Group IV, claim(s) 24, drawn to a method of screening a compound. Election of this group requires an election of species.

Group V, claim(s) 25, drawn to a method of screening a compound for reducing and inhibiting a viral infection.

Group VI, claim(s) 26, drawn to a method of screening a compound for reducing and inhibiting a viral infection using the product of Group I. Election of this group requires an election of species.

Group VII, claim(s) 27, drawn to a method of suppressing a malignant phenotype.

Group VIII, claim(s) 28, drawn to a method of screening a compound for effectiveness in treating a viral infection.

Group IX, claim(s) 29, drawn to a method of screening a compound that can suppress a malignant phenotype in a cell.

The inventions listed as Groups I-IX do not relate to a single general inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, they lack the same or corresponding special technical features for the following reasons:

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In Group I, the special technical feature defining the invention is the nucleotide having specific SEQ ID NOs listed in claim 1. Claims to different products not sharing a structural relationship to the SEQ ID Nos listed, and methods other than the first method of making the using the first product, lack unity of invention.

Group I is drawn to the first product, the nucleic acid sequences listed in claim 1, and the first method of use to reduce or inhibit infection.

Group II is drawn to a second product of a nucleic acid having a regulatory region that does not share the technical feature with Group I.

Group III is drawn to a second method, a method of using a third product, a composition, that inhibits expression or functioning of a gene product.

Group IV is drawn to a third method, a second method of using the first product and a method of using a fourth product, a composition comprising the nucleic acid sequences of Group I.

Group V is drawn to a fourth method, the first use of the second product.

Group VI is drawn to a fifth method, a second use of Group I.

Group VII is drawn to a sixth method, a method of using a fourth product, a composition, which inhibits a malignant phenotype in a cell by inhibiting the functioning of a gene product.

Group VIII is drawn to a seventh method, a method of using a fifth product, a composition, for determining the effectiveness in treating a viral infection.

Group IX is drawn an eighth method, a method of using a sixth product, a compound, that can suppress a malignant phenotype in a cell by increasing the expression level of a gene product.

Groups I-IV and VI contain claims directed to more than one species of the generic invention. These species are deemed to lack unity of invention because they are not so linked as to form a single general inventive concept under PCT Rule 13.1.

Groups I-IV and VI are drawn to nucleotides and nucleotide constructs that contain more than ten individual, independent, and distinct nucleotide sequences in an alternative form.

Accordingly, these claims are subject to restriction under 35 U.S.C. §121 as outlined in 1192

O.G. 68 (November 19, 1996).

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Upon the allowance of a generic claim, applicant will be entitled to consideration of claims to additional species which are written in dependent form or otherwise include all the limitations of an allowed generic claim as provided by 37 CFR 1.141. If claims are added after the election, applicant must indicate which are readable upon the elected species. MPEP § 809.02(a).

The claims are deemed to correspond to the species listed above in the following manner:

Applicant is required to select no more than ten of the individual sequences for examination. The search of no more than ten selected sequences may include the compliments of the selected sequences and, where appropriate, may include subsequences within the selected sequences (e.g., oligomeric probes and/or primers). Alternatively, if applicant wishes to elect a degenerate coding sequence, each of the products recited in claim 5 is seen as a patentably distinct species and election of a single species is required.

Applicant is required, in reply to this action, to elect a single species to which the claims shall be restricted if no generic claim is finally held to be allowable. The reply must also identify the claims readable on the elected species, including any claims subsequently added. An argument that a claim is allowable or that all claims are generic is considered non-responsive unless accompanied by an election.

The following claim(s) are generic: 1, 5, 15, 17, 20, 24, and 26.

The species listed above do not relate to a single inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, the species lack the same or corresponding special technical features for the following reasons: the peptides are different in structure and encode distinctly different nucleic acids that may influence the outcome of the different methods in different ways.

Applicant is reminded that upon the cancellation of claims to a non-elected invention, the inventorship must be amended in compliance with 37 CFR 1.48(b) if one or more of the

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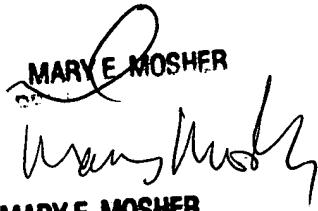
currently named inventors is no longer an inventor of at least one claim remaining in the application. Any amendment of inventorship must be accompanied by a petition under 37 CFR 1.48(b) and by the fee required under 37 CFR 1.17(i).

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Shanon A. Foley whose telephone number is (703) 308-3983. The examiner can normally be reached on 7:30-4:30 M-F.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, James Housel can be reached on (703) 308-4027. The fax phone numbers for the organization where this application or proceeding is assigned are (703) 308-4242 for regular communications and (703) 308-4426 for After Final communications.

Any inquiry of a general nature or relating to the status of this application or proceeding should be directed to the receptionist whose telephone number is (703) 308-0196.

Shanon Foley
January 5, 2001


MARY E. MOSHER
PRIMARY EXAMINER
GROUP 1800
1600

09/509,712

09/509,712

Application No.:

**NOTICE TO COMPLY WITH REQUIREMENTS FOR PATENT APPLICATIONS CONTAINING
NUCLEOTIDE SEQUENCE AND/OR AMINO ACID SEQUENCE DISCLOSURES**

The nucleotide and/or amino acid sequence disclosure contained in this application does not comply with the requirements for such a disclosure as set forth in 37 C.F.R. 1.821 - 1.825 for the following reason(s):

- 1. This application clearly fails to comply with the requirements of 37 C.F.R. 1.821-1.825. Applicant's attention is directed to these regulations, published at 1114 OG 29, May 15, 1990 and at 55 FR 18230, May 1, 1990.
- 2. This application does not contain, as a separate part of the disclosure on paper copy, a "Sequence Listing" as required by 37 C.F.R. 1.821(c).
- 3. A copy of the "Sequence Listing" in computer readable form has not been submitted as required by 37 C.F.R. 1.821(e).
- 4. A copy of the "Sequence Listing" in computer readable form has been submitted. However, the content of the computer readable form does not comply with the requirements of 37 C.F.R. 1.822 and/or 1.823, as indicated on the attached copy of the marked-up "Raw Sequence Listing."
- 5. The computer readable form that has been filed with this application has been found to be damaged and/or unreadable as indicated on the attached CRF Diskette Problem Report. A Substitute computer readable form must be submitted as required by 37 C.F.R. 1.825(d).
- 6. The paper copy of the "Sequence Listing" is not the same as the computer readable form of the "Sequence Listing" as required by 37 C.F.R. 1.821(e).
- 7. Other: _____

Applicant Must Provide:

- An initial or substitute computer readable form (CRF) copy of the "Sequence Listing".
- An initial or substitute paper copy of the "Sequence Listing", as well as an amendment directing its entry into the specification.
- A statement that the content of the paper and computer readable copies are the same and, where applicable, include no new matter, as required by 37 C.F.R. 1.821(e) or 1.821(f) or 1.821(g) or 1.825(b) or 1.825(d).

For questions regarding compliance to these requirements, please contact:

For Rules Interpretation, call (703) 308-4216

For CRF Submission Help, call (703) 308-4212

For PatentIn software help, call (703) 308-6856

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